Page 2

Amendments To the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (currently amended) A compound of the formula I:

I

wherein:

X is selected from the group consisting of:

-NR 10 -, -O-, -CH $_2$ O-, -CONR 10 -, -NR 10 CO-, -CO $_2$ -, -OCO-, -CH $_2$ (NR 10)CO-, -N(COR 10)-, -CH $_2$ N(COR 10)-, phenyl, and C3-6 cycloalkyl,

where R¹⁰ is independently selected from: hydrogen, C₁₋₆ alkyl, benzyl, phenyl, and C₁₋₆ alkyl-C₃₋₆ cycloalkyl, which is unsubstituted or substituted with 1-3 substituents where the substituents are independently selected from: halo, C₁₋₃alkyl, C₁₋₃alkoxy and trifluoromethyl;

W is selected from:

phenyl and heterocycle, which is unsubstituted or substituted with 1-3 substituents where the substituents are independently selected from: halo, C₁₋₃alkoxy and trifluoromethyl;

Z is C;

n is an integer selected from 0, 1, 2, 3 and 4;

Page 3

R1 is selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) trifluoromethoxy,
- (d) hydroxy,
- (e) C₁₋₆alkyl,
- (f) C3-7cycloalkyl,
- (g) -O-C₁-6alkyl,
- (h) -O-C3-7cycloalkyl,
- (i) -SCF₃,
- (j) -S-C₁-6alkyl,
- (k) -SO₂-C₁-6alkyl,
- (l) phenyl,
- (m) heterocycle,
- (n) $-CO_2R^9$,
- (o) -CN,
- (p) $-NR^{9}R^{10}$,
- (q) $-NR^9-SO_2-R^{10}$,
- (r) $-SO_2-NR^9R^{10}$,
- (s) -CONR9R10,
- (t) -NHC(=NH)NR9R10,
- (u) -NHAc,
- (v) $-CH_2C(=O)NHCH_3$,
- (w) $-CH_2C(=O)N(CH_3)_2$,
- (x) -NHCO₂CH₃, and
- (y) hydrogen;
- (t) NHC(=NH)NH², and
- (u) hydrogen,

R9 is selected from H and C1-3alkyl;

R² is selected from:

(C₀₋₆alkyl)-phenyl and (C₀₋₆alkyl)-heterocycle,

where the alkyl is unsubstituted or substituted with 1-7 substituents where the substituents are independently selected from:

(a) halo,

Page 4

- (b) hydroxy,
- (c) -O-C₁-3alkyl,
- (d) trifluoromethyl, and
- (e) -C₁-3alkyl,

and where the phenyl and the heterocycle is unsubstituted or substituted with 1-5 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) trifluoromethoxy,
- (d) hydroxy,
- (e) C₁₋₆alkyl,
- (f) C3-7cycloalkyl,
- (g) -O-C₁-6alkyl,
- (h) -O-C3-7cycloalkyl,
- (i) -SCF₃,
- (j) -S-C₁-6alkyl,
- (k) -SO₂-C₁-6alkyl,
- (l) phenyl,
- (m) heterocycle,
- (n) $-CO_2R^9$,
- (o) -CN,
- (p) $-NR^9R^{10}$,
- (q) $-NR9-SO_2-R10$,
- (r) $-SO_2-NR^9R^{10}$, and
- (s) $-CONR^9R^{10}$;

R³ is -(C₀-6alkyl)-phenyl,

where the alkyl is unsubstituted or substituted with 1-5 substituents where the substituents are independently selected from:

- (a) halo,
- (b) hydroxy,
- (c) -O-C₁-3alkyl, and
- (d) trifluoromethyl,

and where the phenyl is unsubstituted or substituted with 1-5 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,

Page 5

- (c) hydroxy,
- (d) C₁₋₃alkyl,
- (e) -O-C₁₋₃alkyl,
- (f) $-CO_2R^9$,
- (g) -CN,
- (h) -NR9R10, and
- (i) $-CONR^9R^{10}$;

R4 is selected from:

- (a) hydrogen,
- (b) hydroxy,
- (c) C₁₋₆alkyl,
- (d) C₁₋₆alkyl-hydroxy,
- (e) -O-C₁₋₃alkyl,
- (f) $-CO_2R^9$,
- (g) -CONR⁹R¹⁰, and
- (h) -CN;

or where R³ and R⁴ may be joined together to form a ring which is selected from:

- (a) 1H-indene,
- (b) 2,3-dihydro-1H-indene,
- (c) 2,3-dihydro-benzofuran,
- (d) 1,3-dihydro-isobenzofuran,
- (e) 2,3-dihydro-benzothiofuran, and
- (f) 1,3-dihydro-isobenzothiofuran,

or where R^3 and R^5 or R^4 and R^6 may be joined together to form a ring which is phenyl,

wherein the ring is unsubstituted or substituted with 1-7 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) hydroxy,
- (d) C_{1-3} alkyl,
- (e) -O-C₁₋₃alkyl,
- (f) $-CO_2R^9$,
- (g) -CN,
- (h) $-NR^9R^{10}$, and

10/533,337 21056P

Page

6

(i) -CONR9R10; and

R⁵ and R⁶ are independently selected from:

- (a) hydrogen,
- (b) hydroxy,
- (c) C₁₋₆alkyl,
- (d) C₁₋₆alkyl-hydroxy,
- (e) -O-C₁₋₃alkyl,
- (f) oxo, and
- (g) halo;

or a pharmaceutically acceptable salt or individual diastereomer thereof.

2. (previously presented) The compound of Claim 1 of the formula Ia:

Ia

or a pharmaceutically acceptable salt or individual diastereomer thereof.

3. (previously presented) The compound of Claim 1 of the formula Ib:

$$R^3$$
 R^4
 N
 N
 N
 R^2
 R^1

Ιb

or a pharmaceutically acceptable salt or individual diastereomer thereof.

Serial No.:

10/533,337

Case No.: Page 21056P

4. (previously presented) The compound of Claim 1 of the formula Ic:

$$R^7$$
 N
 N
 R^2
 R^3
 N
 R^2
 R^3

Ic

wherein R7 and R8 are independently selected from:

- (a) hydrogen,
- (b) halo,
- (c) trifluoromethyl,
- (d) hydroxy,
- (e) C₁₋₃alkyl,
- (f) -O-C₁-3alkyl,
- (g) -CO₂H,
- (h) -CO₂C₁-3alkyl, and
- (i) -CN;

or a pharmaceutically acceptable salt or individual diastereomer thereof.

5. (previously presented) The compound of Claim 1 of the formula Id:

Id

wherein the dash line represents either single or double bonds;

10/533,337 21056P

Page

8

or a pharmaceutically acceptable salt or individual diastereomer thereof.

6. (previously presented) The compound of Claim 1 of the formula:

$$R^3$$
 R^5
 R^4
 R^6
 N
 N
 R^2
 N
 R^1

wherein W is selected from furanyl, imidazolyl, oxadiazolyl, oxazolyl, phenyl, pyrazolyl, pyrazinyl, pyridyl, pyridazinyl, pyrimidyl, pyrrolyl, thiadiazolyl, and thiazolyl, or a pharmaceutically acceptable salt or individual diastereomer thereof.

- 7. (original) The compound of Claim 1 wherein W is selected from furanyl, imidazolyl, oxadiazolyl, oxazolyl, phenyl, pyrazolyl, pyrazinyl, pyridyl, pyridazinyl, pyrimidyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and triazolyl, and Noxides thereof.
 - 8. (original) The compound of Claim 1 wherein X is -CONH-.
 - 9. (canceled)
 - 10. (previously presented) The compound of Claim 1 wherein n is 0 or 1.
 - 11. (original) The compound of Claim 1 wherein R¹ is selected from:
 - (a) hydrogen
 - (b) halo
 - (c) C_{1-3} alkyl,
 - (d) -O-C₁₋₃alkyl,
 - (e) $-CO_2R^9$,
 - (f) -S-C₁-3alkyl,
 - (g) -SO₂-C₁-3alkyl,
 - (h) -SCF₃,

Page

- (i) $NHC(=NH)NR^9R^{10}$
- (j) $-NR^9R^{10}$,
- (k) $-NR9-SO_2-R_{10}$,
- (l) -SO₂-NR⁹R¹⁰, and
- (m) -CONR⁹R¹⁰.

12. (original) The compound of Claim 1 wherein R² is selected from -(C₀-4alkyl)-phenyl and -(C₀-4alkyl)-heterocycle,

where heterocycle is selected from:

furanyl, imidazolyl, oxadiazolyl, oxazolyl, pyrazolyl, pyrazinyl, pyridyl, pyridazinyl, pyrimidyl, pyrrolyl, thiadiazolyl, thiazolyl, thienyl, and triazolyl, and N-oxides thereof,

where the alkyl is unsubstituted or substituted with 1-7 substituents where the substituents are independently selected from:

- (a) halo,
- (b) hydroxy,
- (c) -O-C₁₋₃alkyl, and
- (d) trifluoromethyl,

and where the phenyl or heterocycle is unsubstituted or substituted with 1-5 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) trifluoromethoxy,
- (d) hydroxy,
- (e) C₁₋₃alkyl,
- (f) -O-C₁₋₃alkyl,
- (g) $-CO_2R^9$,
- (h) -S-C₁₋₃alkyl,
- (i) -SO₂-C₁-3alkyl,
- (j) -SCF₃,
- (k) $-CO_2R^9$,
- (I) $-NR^9R^{10}$
- (m) $-NR^9-SO_2-R^{10}$,
- (n) -SO₂-NR⁹R¹⁰, and
- (o) -CONR9R10.

Page 10

13. (original) The compound of Claim 1 wherein R² is selected from -(C₀-4alkyl)-phenyl and -(C₀-4alkyl)-heterocycle,

where heterocycle is selected from: pyridyl, pyridazinyl, and N-oxides thereof, where the alkyl is unsubstituted or substituted with 1-7 substituents where the substituents are independently selected from:

- (a) halo,
- (b) hydroxy,
- (c) -O-C₁₋₃alkyl, and
- (d) trifluoromethyl,

and where the phenyl or heterocycle is unsubstituted or substituted with 1-3 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) trifluoromethoxy,
- (d) hydroxy,
- (e) C₁₋₃alkyl,
- (f) -O-C₁₋₃alkyl,
- (g) -CO₂-C₁-3alkyl,
- (h) -CO₂H,
- (i) -S-C₁₋₃alkyl,
- (j) -SO₂-C₁-3alkyl,
- (k) -SCF₃,
- (1) -NH₂,
- (m) -NH-SO₂-C₁-3alkyl, and
- (n) -SO₂-NH₂.

14. (original) The compound of Claim 1 wherein R² is selected from -CH₂-phenyl and -CH₂-heterocycle,

where heterocycle is selected from: pyridyl, pyridazinyl, and N-oxides thereof, and where the phenyl or heterocycle is unsubstituted or substituted with 1-3 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) trifluoromethoxy,
- (d) hydroxy,
- (e) C₁₋₃alkyl,

- (f) -O-C₁₋₃alkyl,
- (g) -CO₂-C₁-3alkyl,
- (h) -CO₂H,
- (i) -S-C₁₋₃alkyl,
- (j) -SO₂-C₁-3alkyl,
- (k) -SCF₃,
- (l) -NH₂,
- (m) -NH-SO₂-C₁₋₃alkyl, and
- (n) -SO₂-NH₂.

15. (original) The compound of Claim 1 wherein R² is selected from:

- (1) -CH₂-(phenyl),
- (2) -CH₂-(4-bromophenyl),
- (3) -CH₂-(3-chlorophenyl),
- (4) -CH2-(3,5-difluorophenyl),
- (5) -CH₂-((2-trifluoromethyl)phenyl),
- (6) -CH2-((3-trifluoromethyl)phenyl),
- (7) -CH2-((4-trifluoromethyl)phenyl),
- (8) -CH2-((3-trifluoromethoxy)phenyl),
- (9) -CH2-((3-trifluoromethylthio)phenyl),
- (10) -CH2-((3-trifluoromethoxy-5-thiomethyl)phenyl),
- (11) -CH₂-((3-trifluoromethoxy-5-methoxy)phenyl),
- (12) -CH2-((3-trifluoromethoxy-5-methanesulfonyl)phenyl),
- (13) -CH₂-((3-trifluoromethoxy-5-amino)phenyl),
- (14) -CH2-((3-trifluoromethoxy-5-aminomethanesulfonyl)phenyl),
- (15) -CH2-((3-trifluoromethoxy-5-sulfonylamino)phenyl),
- (16) -CH₂-((3,5-bis-trifluoromethyl)phenyl),
- (17) -CH₂-((3-fluoro-5-trifluoromethyl)phenyl),
- (18) -CH(CH₃)-((3,5-bis-trifluoromethyl)phenyl),
- (19) -C(CH₃)₂-((3,5-bis-trifluoromethyl)phenyl),
- (20) -CH2-(4-(2-trifluoromethyl)pyridyl),
- (21) -CH₂-(5-(3-trifluoromethyl)pyridyl),
- (22) -CH₂-(5-(3-trifluoromethyl)pyridazinyl),
- (23) -CH₂-(4-(2-trifluoromethyl)pyridyl-N-oxide), and
- (24) -CH2-(5-(3-trifluoromethyl)pyridyl-N-oxide).

16. (original) The compound of Claim 1 wherein R³ is hydrogen or phenyl, where the phenyl is unsubstituted or substituted with 1-5 substituents where the substituents are independently selected from:

- (a) halo,
- (b) trifluoromethyl,
- (c) hydroxy,
- (d) C₁₋₃alkyl,
- (e) -O-C₁₋₃alkyl,
- (f) $-CO_2R^9$,
- (g) -CN,
- (h) $-NR^9R^{10}$, and
- (i) $-CONR^9R^{10}$.

17. (original) The compound of Claim 1 wherein R³ is hydrogen or phenyl, where the phenyl is unsubstituted or substituted with 1-3 substituents where the substituents are independently selected from:

- (a) halo,
- (c) hydroxy,
- (d) C₁₋₃alkyl,
- (e) -O-C₁-3alkyl, and
- (f) $-CO_2R^9$.

18. (original) The compound of Claim 1 wherein \mathbb{R}^3 is phenyl, or para-fluorophenyl.

19. (previously presented) The compound of Claim 1 wherein R⁴ is selected from:

- (a) hydrogen,
- (b) hydroxy,
- (c) -CO₂H,
- (d) -CO₂C₁-6alkyl, and
- (e) -CN.

20. (original) The compound of Claim 1 wherein ${\rm R}^5$ and ${\rm R}^6$ are independently selected from:

(a) hydrogen,

(b) hydroxy,

(c) -CH3,

(d) -O-CH3, and

(e) oxo.

21. (canceled)

22. (original) A pharmaceutical composition which comprises an inert carrier and a compound of Claim 1.

23. (canceled)

24. (canceled)

25. (canceled)

26. (original) A method for treating, ameliorating or controlling rheumatoid arthritis which comprises administering to a patient in need thereof an effective amount of the compound of Claim 1.

27. (previously presented) The compound of Claim 1, which is selected from the group consisting of the following compounds, or a pharmaceutically acceptable salt or individual diastereomer thereof:

10/533,337 21056P 14

Ex. 11	O F F F F F F F F F F F F F F F F F F F
Ex. 31	Ex. 32
Ex. 35	Ex. 36
Ex. 37	Ex. 38

10/533,337 21056P

Page

15

N CF3 CF3 CF3	
Ex. 39	
O CF ₃	N CF ₃
Ex. 40	Ex. 41
F O N N N H NH CF ₃	N O CF ₃
Ex. 42	Ex. 43
O N CF ₃	O CF ₃ NHAC
Ex. 44	Ex. 45
N CF ₃ NHCO ₂ Me	O CF ₃ NHSO ₂ Me
Ex. 46	Ex. 47

10/533,337 21056P

Page

16

F O N N H NH CF ₃ NH ₂	P CF ₃ N CF ₃ N N Me H N Me
Ex. 48	Ex. 49
O Me N CF ₃	F O Me CF ₃ NHAc
Ex. 50	Ex. 51
O N N N S NHAC CF ₃ and Ex. 80	ON CF3 NHAC CF3 Ex. 81